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ACCESSION NUMBER:
                          134:57671
DOCUMENT NUMBER:
                         Property for the preparation of 4-alkanoylimidazole
TITLE:
                         derivatives and 1-(2-naphthyl)-1-(1H-imidazol
                         -4-yl alkanol derivatives
                         Kawakamı, Jun-ichi
INVENTOR S.:
PATENT ASSIGNEE(S):
                         Takeda Cherical Industries, Ltd., Japan
                         FCT In. Appl., 39 pp.
SOURCE:
                         CODEN: FIRRE?
                                                     371 BPCT
DOCUMENT TYPE:
                         Fatent
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LANGUACE:
FAMILY ACC. NUM. COUNT:
PATENT INFORMATION:
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     PATENT NO. KIND DATE
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         000078727 A1 230.1238 W0 2000-JP4036 20000621
W: AE, AG, AL, AM, AU, AL, BA, BB, EG, BR, BY, BZ, CA, CN, CR, CU,
     WO 200007872T
             CZ, DM, DZ, EE, GD, GE, HR, HU, ID, N. IN, IS, JP, KG, KR, KZ,
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RW: GH, GM, KE, LS, MW, MZ, SP, SL, SG, TZ, VG, SW, AT, BE, CH, CY, DE, DE, ES, FI, FR, GB, GF, IE, IT, LU, MC, NL, PT, SE, BF, BJ, IF, TG, C1, DM, GA, NN, SW, ML, ME, NE, SN, TD, TG

JP 200104264 A2 21010313 JP 1000-191081 20000631

EP 1193218 A1 21020403 EP 1000-940770 D0100631

R: AT, BS, CH, DE, DK, ES, FE, GB, DE, IT, DI, LU, ND, GE, MC, PT, IE, SI, LT, DV, FI, RO

PRIORITY APELN. IUSO:

JP 1949-175070 A 19490632
WO 20JG-JP4036 W 20000631

OTHER SOURCE(S): CASREACT 134:56671; MARPAT 134:56671

E1 R^7 HO F:6 И Ä NC Ν Γ F.5 R --À. \mathbb{R}^4 F.3 Ι N II F 7 F 1 P.6 \mathbb{R}^2 O V: F:5 F^{\square} F. F 3 F 4 ΙV 11 III

AB An industrially advantageous process for the preph. of compds. of general formula (I; wherein the ring A is an optionally substituted imidazole ring; R is an optionally substituted hydrogarbon group or a heterocyclic group; and E1, E2, E1, E4, E5, R6, and E7 are each hydrogen, optionally substituted hydrogarbyl, OH, SH, NH2, acyl, halogens,

for the like) comprises addn. reaction of 4-cyanoimidazole (II; the ring A is same as above) with F-M1 (E is same as above; M1 = alkali metal, Mg-Y1; Y1 = halo) to give 4-cylimidazole (III; E and ring A are same as above), followed by addn. reaction of EII with haphthalene alkali metals (IV; E1 - E7 are = same as above; M2 is alkali metal, Mg-Y2; Y2 is halo). This process is reduced in the no. of steps, attains a high yield, and dispenses with the use of a heavy metal compd. The compds. I exhibit a steroid C17-G10 lyase inhibitory activity (no data). Thus, a soln. of 42.7 g 4-cyanoimidazole in 50 mL THF was added dropwise to a 1.1. M

scln. of isopropylmagnesium bromide in THF (1.4 L) over a period of 34 min, stirred at 19-25 degree, treated dropwise with $10 \pm$ aq. H2SO4, stirred for

30 min, neutralized to pH 8 with 30 aq. NaOH, and extd. with EtOAc (300 L times. 2) to give 82- 1-(1H-imidazol-4-yl)-2-methyl-1-propanone

(V). 2-Bromo-6-methoxynaphthalene (5.15 g) was added dropwise to a mixt. of 0.55 g and 3 mg iodine in THF at 50.degree. and stirred at 15-25.degree. for 1.5 h, followed by adding dropwise a soln. of 1 g V in THF, and the resulting mixt. was stirred at 15-25.degree. for 8 h to give,

after workup, 84% 1-(1H-imidazol-4-yl)-1-(6-methoxynaphthalen 2-yl)-2-methylpropanol.

REFERENCE COUNT: 19 THERE ARE 19 CITED REFERENCES AVAILABLE FOR THIS

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                  New e-mail delivery for search results now available
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                  FCTFULL has been reloaded
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Jul 14.
                  MOREGE no longer contains STACCARDS file segment
NEWS 11
                  USAN to be reloaded July 28, 2002;
NEWS 13
                  saved answer sets no linger valid
         Jul 19 Enhanced polymer searching in REGISTRY
NEWS 14
         Jul 30 METFIEST to be removed from STM
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NEWS 23
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MEWS EMPRESS February 1 CUFFENT WINDOWS VERSION IS V6.Cd,
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3 Jan 2) Hy 1 G2 Ν G1 4 Ak ² N Н G1 C, H G2 [@1], [@2], [@3], [@4] Structure attributes must be viewed using STN Express query preparation. => s 13 ful FULL SEARCH INITIATED 16:15:34 FILE 'REGISTRY' FULL SCREEN SEAFCH COMPLETED - 49400 TO ITERATE 100.0% PROCESSED 49400 ITERATIONS 4288 ANSWERS SEARCH TIME: 00.00.03 4288 SEA SSS FUL L3 L.1= fil caplus SINCE FILE TOTAL COST IN U.S. DOLLARS **ENTRY** SESSION 280.94 281.15 FULL ESTIMATED COST FILE 'CAPLUS' ENTERED AT 16:15:46 ON 01 OCT 2002 USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT. PLEASE SEE "HELP USAGETERMS" FOR DETAILS. COPYRIGHT (C) 2002 AMERICAN CHEMICAL SOCIETY (ACS) Copyright of the articles to which records in this database refer is hold by the publishers listed in the PUBLISHER (PB) field (available for records published or updated in Chemical Abstracts after December 26, 1996), unless otherwise indicated in the original publications. The CA Lexicon is the copyrighted intellectual property of the American Chemical Society and is provided to assist you in searching databases on STN. Any dissemination, distribution, copying, or storing of this information, without the prior written consent of CAS, is strictly prohibited.

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L8 1514816 INHIBIT?

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desireatives
Kowakano, Jun-Juni
Take by Chemical Industries, Ltd. Japan
POT int. Appl., 35 pp.
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Lipasible INVENTOR(S): PATENT ASSIGNED(S): SOURCE: FOCUMENT TYPE: LANGUAGE FAMIL' ACC. NUM. COUNT: PATENT INFORMATION: عطا سودا APPLICATION NO LATENT HO KING TATE WO 2010078727 A1 0 0 01226 WO 20 W: A0, AG, AL, AM, AU, AU, BA, BA, BB, BG CD, DM, DJ, EF, EJ, GE, HR, HU, ID IL, IN, .S, JP, KS, LC, LK, LP, LT, M, MA, MS, MG, MK, MN, MS, MZ, NO, NZ, PL, PO, FU, SG, SI, JR., 27, TM, TF, TT, UA. 78, UZ. TN, YU, ZA, BT, KG, KJ, MD, RU, TJ, TM RW: GH, GM, KE, LL, MW, MZ, SD, BE, SZ, TZ, UG, LW, AT, BE, CH. CY. DE, PK, ES, Pt. PK, GB, GR, LE, IT. LU. MC, ML, PT, SE, IE, SI, LT, JT, F., RC PRIORITY APPLN INFO:: , LT, JT, F., RC O.: JP 1949-175370 A 19999622 WC 2065-574336 W 2000621 GASRGACT .[4-56872] NARPAT 134:5-871 OTHER SOURCE S:

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LIF ANSWER I OF 7 CAPLUS COPYFIGHT LIGHT ANS CONTINUES.



AB An industrially advantageous **process** for the preph. of compds. of general formula. In wherein the ring A is an optionally substituted.

imidazole ring; F is an optionally substituted hydrocarbon group or a hereroxyclic group; and F1, F2, F3, F4, F4, F6, And F7 are each hydrigen, optionally substituted hydrocarbyl, CH, SH, NH2, acyl, Falogeno, or the like comprises addn. reaction of 4 cyanomimidazole (II; the fire A is sime as above; With P-M1 (F is same as above; M1 = alkali netal, M1, V1, V1 = halo; to give 4-acylimidazile (III; P and ring A are same as another followed by addn. reaction of (II with naphthalene alkali netal) (IV B - R7 are = same as above; M1 is alkali metal, Mg-Y2; Y2 is fail).

The process is reduced in the nol of steps, attairs a high tack, and dispenses with the use of a heavy metal compid. The education of (IV) and dispenses with the use of a heavy metal compid. The corollar is soln of 42.7 q 4-cyanomidazole in 50) mL THF was added dropwise to a . M roln, of isopropylmagnesium bromide in THF (1.4 L) over a [F*Iod of 3) N.n. stirred at 15-25.degree., treated dropwise with 10 aq. 8LYCO4, stirred for 30 min, neutralized to pH 9 with 30 aq NaOH, and exid. M.*1

LIL ANIMER : OF 7 CAPLUS COPYRIGHT 2062 ACS (Continued) RN 30-417-46-1 CAPLUS CN 1-Fippanone, 1-(1H-imidazol-4-yi)- (9CI) (CA INDEX NAME)

H O N C Et N PREFERENCE COUNT: 19 THE

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19 THERE ARE 19 CITED REFERENCES AVAILABLE PECOPD. ALL CITATIONS AVAILABLE IN THE

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LI3 ANSWER 2 OF 7 CAPIUS COPYPLANT 2 2 ACS ACCESSION NUMBER: 2 124 to 2 JAPIUS DOCUMENT NUMBER: 1317315 STILLE: Synthesis, Stilling, and Newtopicte five
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Chainaur, Alaine Crost, Americ Paintaud, Eric,
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AUTHOR S :
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nations superior neuroprotective properties in vivo and in parallel reduced side effects and toxicity. Thus, I administered i.p. protects (8.0)
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              hydroperoxide t BHP) an omidant capable of inducing
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processes. Administration of the archetypal nitrone
phenyl tert-Bu nitrone (PBN) at an equimolar disc also affords some
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intrones were shown to be extremely toxic to rats in contrast to
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                                                                                                                                                                                                                   RN 97744-71-H CAPLUS
UN H-Imidazole 4-markodaldehyae, 7- 4-methoxyphenyl.- (901) - CA
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CN 1H-Imidazole-4-carbonitiale, 2 [4 strifluoromethyl-phenyl] (9CI INDEX NAME) LI3 ANSMER Z OF 7 CAPLUS COPYRISHT. V ACS Continued: RM 279251-02-4 CAPLUS CN IM-Imidazole-4-carboxaldenyde, I Si-naghthalenyl: 4-01 CA IMEEN ARME OHO RN 7/49751-33-5 CAPLUS CN IH Imidazole-4-carbonaldehyde, 2-32 futanyl- 901 °CA INVEX NAME: PN 219251 1446 CARLUT CN UH Imidazole 4-parloxaldebyde, . E furany. 401 TA INCEX NAME: онс н PN 209251 05-7 CAPLUS CN IH Imidazole-4-carboxaldenyde, 7 is threnyl 901. - TA INDEX NAME: H

F. PH . CMSI Re 3 .APTD3 DH .18 Inidazile-4 artonitile,bnlorophenyl . ACC . DA INDEX NAME PH (1975) 474 CARIDS 2M IH Imidazole 4 carbonitrile, 2 3-onlorophenyi 901 COA INDAX NAME H RG 274Lin 98-1 "APIGS CO. 18-imidazole 4-narmonitrile, 1 4 methylphenyl. 901- JPA INTEX NAME. RN = 279251 4)-3 CAPLUS CN = H+ Inidazole 4 Haribonaldehyde, 2-(2-naphthalenyl)= (9C1) = (CA PHOEK NAME) 103 devienemens of H. vs. CH, ...? (@ arbonyl) 113 ANSWER 2 OF 7 CAPLUS COPYRIGHT 2012 ACS H H CHC N RN 279251-07 9 CAPLUS CN 1H-Imidazole 4-carboxaldehyde, 2- 2-thiazolyl) (9CI (CA INSEK NAME) H N N OHC N S 279151 6 . GAPLOS . H Implante 4 carbonaldebyde, 2 of pyriginyl . Well CA INDEX UH NAME N N PH 179251 9-1 PAPIUS CN 1H Imidwayle 4-darbonaldenyde, 2- Brpyridiny. 9-1 FA INCEX NAME SHC H RM = 279251 10 4 CAPTUS CV = 1H Imidazole 4 carboxaldebyde, 2-(4-pyridinyl) + -9015 (CA INTEX NAME. eac N n PH TYPES, TI COMPANY CO. TH. H. Marvie 4 Cast callenges, 7 4 vitightenyl 901 CA INTEX CHARGE

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LI3 ANSWER 3 OF 7
ACCESSION NUMBER:
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INVENTOR OF:

OCHECUTE CONTROL OF THE CONTRO Claude Synthelaho S. A., Fr. U.S., . pp. Conto-in-part of U.S. Ser. No. PATENT ASSIGNEE:S : SOUPOE: 967, .20, 12.694 god date abandoned. CCDEN: USXXAM Patent English LANGUAGE: FAMILY ACC: NUM: COUNT: PATER' INFORMATION: PATENT NO. EIN CATE APPLICATION NO. DATE US 5371277 PE (6835.9 FE 16835.9 US 5432233 PRIORITY APPLIN. INFO.: A. Bī A 1994.2. e 1993 521 1994 211 1995 711 US 1993 3+-.-FR 1991-1:140 1--1.33 US 1394-2954+ FF 1991-1314-M3 1991-1922-2 US 1992-96712 US 1993-39628 1944 -25 19411 28 1441113 1942, 27 OTHER SOURCE(S): MARPAT 123:5587 N p2 AB. The present invention provides a round, which is a quincline derive of the formula Ir RI represents either IH tetrazol-5 yl, or 200H, BZ represents either IH tetrazol-5 yl, or 200H, BZ represents either IV represents either IV represents either IV represents of the represent your representation of each other, hydrogen, nationally approximately 0.1 delevation each other, hydrogen, national style 0.1 delevation expensions of the representation of the repre ea m $_{\rm coll}$ other, by groups of ${\rm CLA}$ along strip, of a CRL bbs group on Which military

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                ● H*1
      10014c 78 4 CALLUS
18-Imidazole 4 carboxaldehyde, C-rutyl b . phenyletneny. — ect
        INCER NAME
Ph. CH. CH. \stackrel{H}{N} Fin
               OH!
      | 150.44-79-5 | CAFIUS | | 18-midazole-4 | arboxaldehyde, 2-bubyl 5- 2-phenylethenyl- ;
on theimicazo:
ethanedicate
(3cI) cA INDEX NAME:
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        CM 2
         CHR 144 €0-7
CMF C2 H2 04
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D13 ANSWER 4 OF 7 CAPIUS COPYRIGHT 2002 ACS ACCESSION NUMBER: 1991:42398 CAPIUS DOCUMENT HUMBER: 134-80 YETTLE:
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1)4:42398

Process for the preparation of Derey biotin
Footsch, Eiker Cisutt, Michael
Merck Fatent Gim bi.H., Fed. Rep. Wer.
U.R., 19 pp. Continuin-part of U.R. 4,877,882.
COLEN: USEXAM
Patent
English
      INVENTOR (S)
        PATENT ASSISNEE (S::
    SOURCE
  DOCUMENT TYPE:
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    NO
        .ANGUAGE
                                                                                                                                                                                                                            English
      LANGUAGE
-AMILY ACC. NUM. COUNT:
PATENT INFORMATION:
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A 1990-626
A1 198011.5
A1 1986-818
A 1986-827
A 1986-827
A 1986-827
A 1987-33
                                              PATENT NO.
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                                                                                                                                                                                                                                                                                                                                                          APPLICATION NO.
US 19-9-422829
TE 199-6-38-0,344
TE 199-7-397-497
US 199-14-16-1
US 199-42-18-8
96
US 493731;

TE (18747

TE (75872)

US 473667

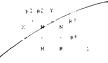
US 483740,

US 487740;

US 487740;

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US 487740;
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    OTHER GOUNCE(S):
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        FI RZ Y
                            N K!
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AB. A procedure for grego, of the title compos, comprises reds, of optically active hydrotos. In Fig. 9.5 H, on substituted alkylicycloadalkylicatalkylichter alkylicotta C, N, or 3/ PIP2 = ion substituted chetero alkylene (conta C, N, or 3/ PIP2 = ion substituted benzyli in Y = 0, ii | II: PP2 = 0 to alms | II: P = H, RF = CH:, etherification to ethers 0, i) III PP+ 10 to alms II P = H. R6 = CH:, etherification to ethers II (P = H: PF = OPH; PS = M. ℓ alkyl), and reaction of the latter with a ryano silane in the presence of a Lewis avid to live distribes II (P = H. R7 = Cyano). Thus, DIBH4 redn. of CVaP \times I (PP+ \times Y = \times PI = Eb, P2 \times H.

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L13 ANSWER 4 OF 7 CAPLUS COPYRIGHT 2002 ACS (Continued) R3 = PhCH2, X = S) gave (7RS,7aR)-1 (R = H, R6 = OH) which was treated by [1,1-carbonyldim.dazole in MeCN to give 'TPS, TaP)-1 (P6 = imidarolyl-1-carbonyloxy). This was treated by Me2SiCN in CH2Cl2 at -20.degree: in presence of TiC.4 to give (TPS, TaPr-1 (Pt = cyano.

Preph. of further D (+) biotin intermediates, e.g.,

/2aS,4S.687-1,3
dibentyl-4--4-carboxyt utyryl tetrahydrothieno(3,4-d)imidarol
-2:3M-one was a.so given.

IT 12938-18-19 112938-18-49 112938-15-59
112938-18-6-69 12938-18-89 112968-29-39
11296-32-89
PD: FCT Feartant,: DRI (Synthetic preparation: FPEP Freparation)
- preph. and reaction of, in preph. of bittin intermediate
PN 12938-13-3 CM-USC
SM SM-Smidaro(1,5-d)intanil tone.
tetrahydro - 5 methowy: caspentyl 3 phenyl++ phenylmethyl (CD) DA INCEX NAME Ph 0 S N N C (CH2)4 OMe Apsolute stereochemistry. Pt. TH2 4



PN 112939-15-5 CARLUS
CN Sthemethiolic alid, S (15-15 methoxy-1 exepentyl)-2-exe 1,3
bis phenylmethyl. 4 imidazolidinyllmethyl, ester. (4P trans--1931) (CA INDEX NAME)

Absolute stereochemistry.

111 ANSWER 4 OF T CAPIDS TOFFFISHT 2 2 A S. T. Settleman Ph N 3MH 18, 4 3MH FN 11 939 F F TAILUS 18 4 United Dissipation of the Company of th Abbelote stereochemistry. PE o N CH₂ 4 CC₂H PN 11998 to 4 CARLON
CN 18 3B Imagazo(1,5%, thiazole-7 massicualdebyte,
tersanydro-5-oxo a phenyl r
-phenylmethyly 201 JA INCEN NAME) CH2 Ph 0 BN 117568-23-3 CAPIDS CN 18-38 Impdato[1,5-d]thrapple-T-carbinitrie, terrah dro 5-oxo 3-phenyl -phenylmethy. - . . - CA IMIEX NAME . CH2 Ph ٤ 0

L1. ADDRESS OF 7 CAPLUS COPYRIGHT 2012 ACS
ACCESSION NUMBER:
DOCUMENT NUMBER:
TITLE:
INTENTION:S.:
PROMES for the preparation of 1-(+)-biotin
PROMES AUSIGNEE(S):
SOUFCE
DOCUMENT TYPE:
DOCUMENT TYPE:
LANCUAGE
FAMILITY ACC. NUM. COUNT:
FAMILITY ACC. NUM. COUNT:
DETENT LIFORMATION:

TO CAPTURE 2012 ACS
ACCESSION NUM. ACCESS
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THE COPYRIGHT 2012 ACS
ACCESSION NUMBER:
THE COPYRIGHT 2012 ACCESSION NUMBER:
THE COPYRIGHT 20 DOCUMENT TYPE: LANGUAGE FAMILY ACC. NUM. COUNT: PATENT INFORMATION: PATENT NO. KIND DATE 14991.31 18911.5 1483.816 1483.827 1484.827 1484.626 1447.8315 PRICE TO APPLING THE PRICE THE

OTHER SIUPPE S : MARIAT 113:23:41

2 pl p2 7 PROHEN NORIPE K N NP 3 S (CH2) 4CO(H 111 . p4 11 64

AB The process for project the title sompore I is characterized in that the synthesis is carried out via an intermediate II FL F2 of

H, Substituted askyl, mysloackyl, asyl, asylly, betestastyl wherein the heteroatom is N, 7, 3; PIPC # surstitutes ackylete. heteroatkylene: P3 # isabstituted thrHI; PAPF = D1 X, Y = 1, X | ty sir ensite reds. to

an alo., esterification. E oversion to nitrile, to an alkanoyi, cleavage to

imidatolidine and y liming to a I deriv. worsh by known methods
is nonverted to I, or conversion of the alkanoyl with an adult to a

113 ANSWER 4 OF 1 PARLIES of FIREIGHT . | LACE | Lacetower

PH 112968 30 8 CAPING ON 1H, 3H Immidstyll, 50 1H, 3H Immidstyll, 50 citizatile 3 hexandic wold, tetrahydro-epsilon. 5 dilko 3 prenyl c şlenyimetnyl 501 504 IMMEX DAME

Er. 0 S N N

C 38, 4 11, 8

LIS ANSWER 5 OF 7 CAPLUS COPERIGHT 2002 ACS "Continued derivs, which by known methods is converted to 1.

23.63, 485, 688::III was prepd.

IT 112938-13-3P 112938-14-4P 112938-15-5P 112938-13-3P 112938-13-5P 112938-13-3P 112948-32-8P RI: RCT (Reactant), SPN (Synthetic preparation): PPEP (Preparation) (prepn. and reaction of, in prepn. of biolin intermediate)

RN 112948-13-3 CAPLUS

CN 34, 584-1midato(1,5 c)thiazol-5-one, tetrahylarol-5-senehoxy: coopenty1-3-pheny1-6-(phenylmethy): 901 CA INCEX NAME Fh 0 S N N C :CH2 4 PMe 0 PH 112939-14 4 CAPLUS
TH 2-imidatolidinone,
4 meloaptomethy: 5 -5 methoxy-loxopentyl (1,)
his-phenylmethyl , 49 trans (PCI TA IMIEX NAME

Atsolute stereschemistry

Ph S N R CH2 4

PG | 112988-15-5 CAPLUS

VN | Sthemethiolic alis. B | [5 | f methoxy | 1 exopenty, | 7 | 20 | 1,3tisiphenylmethy) | 4 imidatolidinyl[methyl, | ester. | 48 trans
Sci | [56] | INDEX NAME

Absolute stereconemistry

113 ANSWER S DE COMPINS CONTRETED LOS ANA CONTRIBUTA

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THE STATE OF THE STATE

S N

PN 117'08 PM 3 CAPEDO CN 1H, BH Imstaco[1,5 officiarole foarDunitries, tetranydro 5 oxo 3 phenyl (spenylmethyl) (971) CA IMER WAME

CH2 Ph N 0

Ph O S N N с тенути соун

LIB ANSWER 6 OF 7 CAPLUS COPYRIGHT 2 - 2 ACS
ACCESSION NUMBER: 1988:112-77 CAPLUS
EOCUMENT NUMBER: 1988:112-77 CAPLUS
TITLE: 1981-12-77 CAPLUS
FACENT ASSINER(ST): 5 Process for the preparation of 1 - Eightn
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PATENT NG. FIND TATE APELIDATION NO. 1ATE BE 14811514 1441406 EF 142686 A7 198 100 d EF 142686 A3 198 100 d EF 142686 B1 198 010 EF 142681 B1 198 010 EF 142681 B1 198 010 EF 157 188 A1 198 414 PPIOPITY APPLING INEXCO 18 1-- 3813/45 1996 419 18 2997 35 3572 1997 2 5 18 1996 561245 2556 419 18 1997 37 38 2 1997 2 5

p1 p2 V N N NF A . p4

A procedure for the preph. of 1 - flotin from 1 systeme or

L serine via hisystic intermediate I [RI.R] - H. unsubstitited alkyl, pyloalkyl, aryl, aralkyl, heteroaryl; FiF. In Kinstituted alkyler.

heteroalkylene: Ps \in H, N-grote-tive group. F4R5 = 0: X, Y = 0, S}.

is described DiBM4 redn. of TH 1 P1 Ph. PT H. As 1 En HZ. R4P1 T Y =

1.3 ANOMER SIGE COMPANY CONTENTS OF ACCUSE OF THE PROPERTY OF

:T 112938-18-8P IT 11938-18-8P

RIF ROT ORGANISHED (SYMICHYMETER) Preparation , PREP Treparation of the file day of the man O acetylation of the file day of the man O acetylation of the file day of the man O acetylation of the file day of the man O acetylation of the file day of the fi

CH9 PH2 IN N c

Fh. 5 SH2 PH C CH2 4 SMe 6

Et 1119:9-14 4 MARION MO 2-importalistiche, 4 merzeignetty, 10 5 delnowy 1 oxigenty, 103 EX NAME 118 (henylethy) / 48 frank MO DA INTEX NAME

Adsolite stereotherustry.

0 3 B B ZHD 4 OME

EN 112939 Ust CAPLUS

ON Ethanethiolicavid, 20035 t methody coopenty. I oxil, 8
tis prenylmethy. 4 imidazolidiny.[methyl, ester. 45 trans 401 28
INCEX 104ME

Assiste stereothemistry

1.3 ANAMER + DE 7 CARTON NIGHT-HILL NAV. (mg/intinged

Es N P SAC

Ansolité stereochemistry.



PN 11939 1946 CAPLUS CN HK.9H Imidazo[1.5] (Initazole ' harboxaldenyon, tetrahydro 5 oko-3 phenyl ri (phenylmenhyl) (901) (MA INDEX MAME)

CH₂ Fh THO 5 54

RN 112968 T9 3 CAPLUS CN 1H, 3H-Imidazo(1, 5-o)thrazole 7 darbouitrile, tetrahydro 5 oxo-3-phenyl-+ (prenylmethyl) 9-011 JA INCEX NAME.

L13 ANSWER 7 OF 7 CAPLUS COPYRIGHT 2002 ACS ACCESSION NUMBER: .982:4-6222 CAPLUS

US COPPTION LUC ACS
,981:44-6222 CAPUNS
,97:6-222
Synthesis of 4,5-disubstituted imidazoles
Kavadias, Gerry Luh, Bing: Saintonge, Roger
Bristel Lab. Canada, Canadas, Pg. 158-171, Can.
Can. J. Chem. 1992 , 6-16-, 723-9
COLEM: LICHAG: ISSN: 7.78-4-42
Dournal DOCUMENT NUMBER: TITLE: ALTHOR(S): CORPORATE SOURCE: SCUPCE:

DOCUMENT TUPE: LANGUAGE: OTHER BOURGE'S': GI English CASPEACT 97:6222

CHC KIH, DH CH) H HC:(H)

AB Introduction of Ph}C on the N of imidazole 4,5 dicarboxyliv and estera or 4,5 dihydroxymethylimidazole deactivated the functional

group
as acent to the protecting group and allowed seartions to take prace preferably or exclusively on the other fun timbal group. Thus, di-Me

. I triphenylmethylimidabole 4, dibarboxylate I , on treatment with NHINE

and MeNH2 produced Me

and MeNH: produced Me
4-nydrad.nocationyl-intriphenylmethylimidatole 5
carboxylate and Me
4-methylam nocationyl-intriphenylmethylimidatole 5
carboxylate, resp. Pedn. of 1 with LiBH4 gave Me
4-hydroxymethyl :
t typhenylmethylimidatole 5 darroxylate. Treatment of 4.5b sshydroxymethyl of triphenylmethylimidatole 11 , with Mei. No.
and with

With Augo afforded 5-bydronymetny, 4 trimetnyla etikymetnyl-1-triphenylmetnylimidabole, and 4 arehoxymetnyl 1 hydroxymetnyl 1 triphenylmethylimidabole, resp. Ixido of 11 with activated Modd

produced

the monoaldehydes III and IV in a ratio of IV II. A new mild process for deprotection of N triphenylmethylimidateles, compatible with and sensitive groups in the mol , is reported. sunthesis of several 4.5 insufstituted imidazoles is also

described. 1T 82032-50-6P .. egusarourem Plo EPE ofroperties of EEG Cymonetic preparation of EMEE Preparation prepulance 1999 of

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Ph 1 THO Po s t N с сні 4 кори

113 ANSWER 7 OF 7 CAPLUS COPYRIGHT 2002 ACS (**Continued) RN 82030-50-6 CAPLUS (**Continued) RN 82030-50-6 CAPLUS (**Continued) RH Emidazole-5-cailoxaldehyde, 4-(hydroxymethyl)-1-(triphenylmethyl)-501 (**CAINDEX NAME)

N CH2 OH N

82032-51-7P

ceb.

N CH; OH

8

TA INDEX NAME

● #51

PN HOSE OF CHECK NAME

ON THE Inidacie 4 Paramitrile, I hydroxymethyl , monohydromic, he set on the NAME

ON CHECK NAME

inget redevence

Vs. claim 17 Por 10/019094

ACCESSION NUMBER: EGCUMENT NUMBER:

TITLE:

24 ANSWER 18 OF 18 CAPLUS COPYRIGHT 2002 ACS 1977:106471 CAPLUS

36:106471

Photochemical reactions. Part 91. Photochemistry of imidazolides. I. The photo-Fries-type rearrangement

of N-substituted imidazoles

Iwasaki, Shigeo AUTEOR(ε): CORPORATE SOURCE:

Org.-Them. Lab., ETH, Zurich, Switz. 👍 Helv. Chim. Acta (1976), 59(8), 2733-52

CODEN: HCACAV

Journal Er.alish

IDOUMENT TYPE:

LANGUAGE:

SOURCE:

GI

Ν ΙI III

Imidazoles I (R = Ac, Me(CH2)6CO, cyclohexylcarbonyl, Me3CCO, Bz, Me2C:CHCO, MeO2C, Et2NCO, PhCH2) underwent photochem. rearrangements to give 6-45% II and 10-35% III. The structures of II and III were confirmed

by spectral data, which are reported.

61985-31-7P ΙT

F.L: PREF (Preparation)

(by photochem, rearrangement of 1-acyl analog)

61385-31-7 CAPLUS

1-Fropanone, 1-(1H-imidazcl-4-yl)-2,2-dimethyl- (9CI) (CA INDEX NAME) CII

